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(54) Title: FRAGRANCE COMPOSITIONS

(57) **Abstract:** The present invention relates to a composition comprising: (a) a fragrance oil wherein the fragrance oil comprises: (i) one or more perfume raw materials with a high odour impact which have an odour detection threshold of less than, or equal to about 50 parts per billion; (ii) less than about 5 %, by weight of the fragrance oil, of top note perfume raw materials where in the top note perfume raw materials have a boiling point of less than 250 °C at 1 atmosphere pressure; (b) an entrapment material which is selected from the group consisting of polymers; capsules, microcapsules, and nanocapsules; liposomes; film formers; absorbents; cyclic oligosaccharides and mixtures thereof; wherein the perfume raw material and the entrapment material exist in an associated form on the substrate and wherein the weight ratio of high odour impact perfume raw materials which have an odour detection threshold of less than, or equal to, about 50 parts per billion to entrapment material within the associated form falls in the range of about 1:20 to about 20:1. The present invention provides compositions wherein the fragrance character remains detectable for greater than about 2 hours, preferably greater than about 4 hours, more preferably greater than about 6 hours, after the composition has been applied to the substrate.

Fragrance Compositions

Field of the Invention

The present invention relates to compositions, in particular cosmetic compositions, more particularly fragrance compositions, which comprise a fragrance oil and an entrapment material, wherein the fragrance oil comprises a low level of "top note" perfume raw materials and a high level of "high odour impact" perfume raw materials. More particularly this invention relates to compositions wherein the fragrance character of the pleasant "high odour impact" perfume raw materials remains detectable on the substrate for at least 2 hours after application. Compositions of the present invention are suitable for application to a wide variety of substrates but particularly to the skin and hair.

Background to the Invention

It has long been a feature of many types of compositions, including cosmetic compositions, that they comprise a fragrance oil for the purpose of delivering a pleasant smell. This can improve the overall consumer acceptance of the composition or mask unpleasant odours. In fact, it can be the sole purpose of some compositions to impart a pleasant odour to suitable substrates including the skin and or hair.

Fragrance oils used within such compositions usually comprise many different perfume raw materials. Each perfume raw material used differs from another by several important properties including individual character, and volatility. By bearing in mind these different properties, and others, the perfume raw material can be blended to develop a

fragrance oil with an overall specific character profile. To date, characters are designed to alter and develop with time as the different perfume raw materials evaporate from the substrate and are detected by the user. For example perfume raw materials which have a high volatility and low substantivity are commonly used to give an initial burst of characters such as light, fresh, fruity, citrus, green or delicate floral to the fragrance oil which are detected soon after application. Such materials are commonly referred to in the field of fragrances as "top notes". By way of a contrast, the less volatile, and more substantive, perfume raw materials are typically used to give characters such as musk, sweet, balsamic, spicy, woody or heavy floral to the fragrance oil which, although may also be detected soon after application, also last for longer. These materials are commonly referred to as "middle notes" or "base notes". Highly skilled perfumers are usually employed to carefully blend perfume raw materials so that the resultant fragrance oils have the desired overall fragrance character profile. The desired overall character is dependent both upon the type of composition in which the fragrance oil will finally be used and also the consumer preference for a fragrance.

In addition to the volatility, another important characteristic of a perfume raw material is its olfactory detection level, otherwise known as the odour detection threshold. If a perfume raw material has a low odour detection threshold, otherwise known as "high odour impact", only very low levels are required of the raw material within a composition for it to be detected by the user, sometimes as low as a few parts per billion. Conversely, if a perfume raw material has a high odour detection threshold, otherwise known as a "low odour impact", larger amounts of that material are required before it can be smelt by the user. To date, when developing a fragrance oil, it has been important to balance the fragrance with both low and high odour impact perfume raw materials since the use of too many high odour impact materials could lead to a short lived overwhelming scent. As such the levels of high odour impact perfume raw materials within a fragrance oil have traditionally been restricted.

To date, due to the physical characteristics of the perfume raw materials, ie the volatility and the odour detection impact, the possible types of overall characters of fragrances have

been limited. In addition, it has been difficult to develop a fragrance oil which remains detectable on the substrate over very long periods of time. Furthermore, it has not always been possible to develop a fragrance character which retains its initial fragrance character over time. However, it is known that consumer preference for fragranced compositions is mostly driven by the initial character detected after application. It is therefore desirable to have a fragrance which retains its initial character over time, and wherein that character remains noticeable to the consumer over time. Since materials with high odour impact can be detected when only very small levels are present, they are particularly useful for developing a noticeable long lasting fragrance character. It is also desirable to be able to create new to the world fragrance character profiles wherein one, or several, well recognised characters, regardless of their volatility, are maintained overtime such that a unique long lasting, "top, middle and base note" character is created. As such, it would be advantageous to be able to create a fragrance oil which will impart, in a new way, fragrance materials with a high odour impact, such that they can be experienced both initially, and over a substantial period of time, by the user.

In the past, many attempts have been made to delay the volatility profiles of fragrance oils within many types of compositions to extend the overall fragrance effect. For instance the fragrance oil may be formulated to include a higher proportion of perfume raw materials with a low volatility and which are therefore more substantive on the substrate. However, as discussed above, this restricts the fragrance character that can be achieved over time. Another approach has been to chemically, and reversibly, modify the perfume raw materials to a pro-perfume compound which is disclosed in patent applications WO 98/47477; WO 99/43667; WO 98/07405; WO 98/47478; all of which are incorporated herein by reference. The resultant pro-perfumes are not themselves volatile but, after the chemical modification is reversed, usually by hydrolysis upon application to the substrate, the perfume raw material is released and can then evaporate in the usual way. In these examples the release rate of the perfume raw materials is controlled by the reaction rate of the pro-perfume to perfume raw material transformation.

Further disclosures have discussed improving the overall longevity of a fragrance by delaying the evaporation of the fragrance oils. A wide variety of techniques have been disclosed among them encapsulation of the perfume raw materials for example within capsules (disclosed in JP-A-58/052211, EP-A-303,461), absorbing the materials to a surface for example by using carbon or zeolites (disclosed in US 6,033,679), occluding the release of the perfume raw materials for example by the formation of a film (disclosed in US 3,939,099) and complexing the perfume raw materials for example by using cyclic oligosaccharides. The prior art on this latter method includes JP-A-6/287127 and JP-A-8/176587 which disclose use of hydroxyalkylated cyclodextrins within cosmetic compositions to sustain the effect of the fragrance; and JP-A-8/183719 and JP-A-10/120541 which discloses a combination of cyclodextrin encapsulated fragrance and non encapsulated fragrance within a deodorant composition for prolonging the fragrance duration to at least 2 hours, all of which are incorporated herein by reference.

Whilst the compositions and disclosures of the prior art provide useful teachings for prolonging the overall fragrance character of different compositions the approaches still have limitations. The pro-perfume approach is limited by those chemical modifications that can suitably be made to the perfume raw materials. In addition, the prior art is restricted by pro chemistries used, or by the use of low levels of pro perfumes, thus preventing the development of a fragrance oil which exhibits a broad range of top note characters over time. On the other hand, entrapment materials, when used in the traditional way, interact with a broad range of perfume materials including top, middle and base notes prolonging the overall character of the whole fragrance. As such the prior art does not sufficiently teach how to preferentially delay the evaporation of a large range of high odour impact perfume raw materials within a single fragranced composition such that these distinct characters can be recognised throughout the usage period. In addition, since, in general, entrapment such as that described suppresses the evaporation of only a relatively small amount of the perfume raw materials, the low level of delayed release is often not noticeable to the user. As such the prior art does not adequately teach how to provide a fragrance wherein the prolonged character has sufficient strength to be noticeable and remains distinctive over time.

Surprisingly, it has now been found that compositions comprising a fragrance oil and a material which is able to delay the evaporation of the fragrance oil, wherein the fragrance oil is blended to comprise low levels of "top note" perfume raw materials and wherein the weight ratio of "high odour impact" perfume raw materials which have an odour detection threshold of less than, or equal to, 50 parts per billion to entrapment material within the associated form falls in the range of 1:20 to 20:1, can be used to create a long lasting fragrance character profile which has prolonged, and noticeable distinct characters. In addition, it has been found that within such fragrance oils, much higher levels of "high odour impact" perfume raw materials can be used than would be acceptable in a traditionally developed fragrance which leads to new and distinct fragrance characters. Furthermore, this use of high levels of "high odour impact" perfume raw materials can result in a reduction in the level of fragrance oil that is required in the final finished product which can reduce skin sensitisation and can have cost reduction benefits both for the manufacturer and the consumer. By controlling the weight ratio between the entrapment material and the "high odour impact" perfume raw materials highly efficient use is made of the entrapment material which can also lead to a reduction in the amount of both fragrance oil and entrapment material required in any given formulation. In addition, this use of "high odour impact" perfume raw materials can also result in a reduction in the level of entrapment material required to achieve the desired benefit which can have both aesthetic benefits for the final formulation in addition to further cost saving benefits.

While not wishing to be bound by theory, it is believed that when the entrapment material is reversibly associated to the perfume raw materials that the compositions of the present invention are substantially free of fragrance - entrapment material complexes until applied to the substrate. Upon application the volatile solvent evaporates providing an initial fragrance burst while the remaining fragrance complexes with the entrapment material. Over time, this association breaks down resulting in release of the perfume raw materials. Since the composition comprises a fragrance oil which has been developed with a high level of "high odour impact" perfume raw materials, the very low levels of materials that

are released from the association over time are easily detected by the user. In addition, despite the fact that the fragrance oil is developed with very high levels of "high odour impact" perfume raw materials, the association that forms means that the fragrance is not overwhelming when it is first applied.

It is an object of the present invention to provide compositions, which impart a long lasting and noticeable fragrance character to the substrate on which they are applied. It is a further object of this invention to provide a fragrance which maintains its initial character profile over time in a noticeable way. This, and other objects of this invention, will become apparent in light of the following disclosure.

Summary of the Invention

The present invention relates to a composition comprising:

- (a) a fragrance oil wherein the fragrance oil comprises:
 - (i) one or more perfume raw materials with a high odour impact which have an odour detection threshold of less than, or equal to about 50 parts per billion
 - (ii) less than about 5%, by weight of the fragrance oil, of top note perfume raw materials where in the top note perfume raw materials have a boiling point of less than 250°C at 1 atmosphere pressure
 - (b) an entrapment material which is selected from the group consisting of polymers; capsules, microcapsules, and nanocapsules; liposomes; film formers; absorbents; cyclic oligosaccharides and mixtures thereof;
- wherein the perfume raw material and the entrapment material exist in an associated form on the substrate and wherein the weight ratio of high odour impact perfume raw materials which have an odour detection threshold of less than, or equal to, about 50 parts per billion to entrapment material within the associated form falls in the range of about 1:20 to about 20:1.

This invention further relates to methods of use for such compositions and the use of such compositions to delay the release of the perfume raw materials from the substrate.

Detailed Description of the Invention

All percentages and ratios used herein are by weight of the total composition and all measurements made are at 25°C, unless otherwise designated. Unless otherwise indicated all percentages, ratios and levels of ingredients referred to herein are based on the actual amount of the ingredient, and do not include solvent, fillers or other materials which may be combined with the ingredient in commercially available products.

All publications cited herein are hereby incorporated by reference in their entirety, unless otherwise indicated.

The term "dermatologically-acceptable," as used herein, means that the compositions, or components thereof, are suitable for use in contact with human skin without undue toxicity, incompatibility, instability, allergic response, and the like.

The term "safe and effective amount" as used herein means an amount of a compound, component, or composition sufficient to significantly induce a positive benefit, but low enough to avoid serious side effects, i.e. to provide a reasonable benefit to risk ratio, within the scope of sound medical judgement.

Active and other ingredients useful herein may be categorised or described herein by their cosmetic and/or therapeutic benefit or their postulated mode of action. However, it is to be understood that the active and other ingredients useful herein can in some instances provide more than one cosmetic and/or therapeutic benefit or operate via more than one mode of action. Therefore, classifications herein are made for the sake of convenience

and are not intended to limit an ingredient to the particularly stated application or applications listed.

The elements of these compositions are described in more detail below.

Fragrance Oil

Compositions of the present invention comprise from about 0.01% to about 99%, preferably from about 0.25% to about 50%, more preferably from about 0.5% to about 40%, even more preferably from about 1% to about 25%, and most preferably from about 2.5% to about 25%, by weight, of the fragrance oil.

As used herein the term "fragrance oil" relates to a perfume raw material, or mixture of perfume raw materials, that are used to impart an overall pleasant odour profile to a cosmetic composition. In addition "perfume raw materials" are defined by materials with a ClogP value preferably greater than about 0.1, more preferably greater than about 0.5, even more preferably greater than about 1.0. As used herein the term "ClogP" means the logarithm to base 10 of the octanol / water partition coefficient. This can be readily calculated from a programme called "CLOGP" which is available from Daylight Chemical Information Systems Inc., Irvine CA, USA. Octanol / water partition coefficients are described in more detail in US-A-5,578,563.

Within the present invention, by mixing together several different perfume raw materials a fragrance oil can be achieved which, when used in a composition in conjunction with an entrapment material, is able to impart a particular long lasting character to the composition in which it is used. The mixture of perfume raw materials used will be carefully chosen and blended to achieve a fragrance oil with the desired overall fragrance character profile.

The fragrance oil itself can comprise any perfume raw material suitable for use in compositions. Overall the fragrance oil will most often be liquid at ambient temperatures and consist of a single individual perfume raw material. A wide variety of chemicals are

known for fragrance uses, including materials such as aldehydes, ketones and esters. However, naturally occurring plant and animal oils and exudates comprising complex mixtures of various chemical components are also commonly known for use as fragrances. The individual perfume raw materials which comprise a known natural oil can be found by reference to Journals commonly used by those skilled in the art such as "Perfume and Flavourist" or "Journal of Essential Oil Research". In addition some perfume raw materials are supplied by the fragrance houses as mixtures in the form of proprietary speciality accords.

In order that fragrance oils can be developed with the appropriate character for the present invention the perfume raw materials have been classified based upon two key physical characteristics:

- (i) boiling point (BP) measured at 1 atmosphere pressure. The boiling point of many fragrance materials are given in *Perfume and Flavor Chemicals (Aroma Chemicals)*, *Steffen Arctander (1969)*. Perfume raw materials for use in the present invention are divided into volatile raw materials (which have a boiling point of less than, or equal to, about 250°C) and residual raw materials (which have a boiling point of greater than about 250°C, preferably greater than about 275°C). Volatile raw materials, for the purposes of this invention, are considered to be those that impart "top note" ie light, fresh, fruity, citrus, green or delicate floral characters to the fragrance oil and the like. Similarly the residual perfume raw materials are considered to be those that impart "middle or base note" ie musk, sweet, balsamic, spicy, woody or heavy floral characters to the fragrance oil and the like. All perfume raw materials will preferably have boiling points (BP) of about 500°C or lower.
- (ii) odour detection threshold which is defined as the lowest vapour concentration of that material which can be olfactorily detected. The odour detection threshold and some odour detection threshold values are discussed in eg "Standardized Human Olfactory Thresholds", M. Devos et al, IRL Press at Oxford University Press, 1990, and "Compilation of Odor and Taste Threshold Values Data", F. A. Fazzalar, editor ASTM Data Series DS 48A, American Society for Testing and Materials, 1978, both of said publications being incorporated by reference. Perfume raw materials for use in the

present invention can be classified as those with a low odour detection (ie "high odour impact") with a detection threshold of less than 50 parts per billion, preferably less than 10 parts per billion and those with a high odour detection (ie "low odour impact") with a threshold of greater than 50 parts per billion (values as determined from the references above).

Since, in general, perfume raw materials refer to a single individual compound, their physical properties (such ClogP, boiling point, odour detection threshold) can be found by referencing the texts cited above. In the case that the perfume raw material is a natural oil, which comprises a mixture of several compounds, the physical properties of the complete oil should be taken as the weighted average of the individual components. In the case that the perfume raw material is a proprietary speciality accord the physical properties should be obtain from the Supplier.

In order to develop fragrance oils that are suitable for use in the present invention it is necessary that the fragrance oil comprises less than about 5%, preferably less than about 4%, more preferably less than about 3% and even more preferably less than about 2%, by weight of the fragrance oil, of top note perfume raw materials. In addition the fragrance oil should comprise sufficient levels of "high odour impact" perfume raw materials such that the weight ratio of high odour impact perfume raw materials which have an odour detection threshold of less than, or equal to, about 50 parts per billion to entrapment material within the associated form falls in the range of about 1:20 to about 20:1, preferably from about 1:10 to about 10:1, more preferably from about 1:7 to about 7:1, even more preferably from about 1:5 to about 5:1, further more preferably from about 1:3 to about 3:1 and most preferably from about 1:2 to about 2:1.

It is preferred that the fragrance oils for use herein comprise from about 0.1% to about 99%, preferably from about 1% to about 75%, more preferably from about 5% to about 50%, even more preferably from about 20% to about 40%, by weight of the fragrance oil, of "high odour impact" perfume raw materials which have an odour detection threshold of less than, or equal to, about 50 parts per billion. It is preferred that the high odour impact

perfume raw materials have an odour detection threshold of less than, or equal to about 40 parts per billion, preferably about 30 parts per billion, more preferably about 20 parts per billion and even more preferably about 10 parts per billion.

It is a further feature of the present invention that, since the fragrance oil comprises a high level of perfume raw materials with a high odour impact, the compositions can comprise lower levels of fragrance oil than would traditionally be present. As such, compositions of the present invention can preferably comprise from about 1% to about 15%, preferably from about 1% to about 8%, and more preferably from about 2% to about 6%, by weight, of fragrance oil.

In general a broad range of suitable perfume raw materials can be found in US-A-4,145,184, US-A-4,209,417, US-A-4,515,705, and US-A-4,152,272. Non-limiting examples of perfume raw materials which are useful for blending to formulate fragrance oils for the present invention are given below. Any perfume raw materials, natural oils or proprietary speciality accords known to a person skilled in the art can be used within the present invention.

Examples of residual "middle and base note" perfume raw materials having a boiling point of greater than 250°C, which have a low odour detection threshold are selected from, but are not limited to, ethyl methyl phenyl glycidate, ethyl vanillin, heliotropin, indol, methyl anthranilate, vanillin, amyl salicylate, coumarin. Further examples of residual perfume raw materials having a boiling point of greater than 250°C which are generally known to have a low odour detection threshold include, but are not limited to, ambrox, bacdanol, benzyl salicylate, butyl anthranilate, cetalox, ebanol, cis-3-hexenyl salicylate, lilial, gamma undecalactone, gamma dodecalactone, gamma decalactone, calone, cymal, dihydro iso jasmonate, iso eugenol, lyrat, methyl beta naphthyl ketone, beta naphthol methyl ether, para hydroxyl phenyl butanone, 8-cyclohexadecen-1-one, oxocyclohexadecen-2-one / habanolide, florhydral, intreleven aldehyde.

Examples of volatile "top note" perfume raw materials having a boiling point of less than, or equal to, 250°C, with a low odour detection threshold are selected from, but are not limited to, anethol, methyl heptine carbonate, ethyl aceto acetate, para cymene, nerol, decyl aldehyde, para cresol, methyl phenyl carbonyl acetate, ionone alpha, ionone beta, undecylenic aldehyde, undecyl aldehyde, 2,6-nonadienal, nonyl aldehyde, octyl aldehyde. Further examples of volatile perfume raw materials having a boiling point of less than, or equal to, 250°C, which are generally known to have a low odour detection threshold include, but are not limited to, phenyl acetaldehyde, anisic aldehyde, benzyl acetone, ethyl-2-methyl butyrate, damascenone, damascone alpha, damascone beta, flor acetate, frutene, fructone, herbavert, iso cyclo citral, methyl isobutenyl tetrahydro pyran, iso propyl quinoline, 2,6-nonadien-1-ol, 2-methoxy-3-(2-methylpropyl)- pyrazine, methyl octine carbonate, tridecene-2-nitrile, allyl amyl glycolate, cyclogalbanate, cyclal C, melonal, gamma nonalactone, cis 1,3-oxathiane-2-methyl-4-propyl.

Other residual "middle and base note" perfume raw materials having a boiling point of greater than 250°C which are useful in the present invention, but which have a high odour detection threshold, are selected from, but are not limited to, eugenol, amyl cinnamic aldehyde, hexyl cinnamic aldehyde, hexyl salicylate, methyl dihydro jasmonate, sandalore, veloutone, undecavertol, exaltolide/cyclopentadecanolide, zingerone, methyl cedrylone, sandela, dimethyl benzyl carbonyl butyrate, dimethyl benzyl carbonyl isobutyrate, triethyl citrate, cashmeran, phenoxy ethyl isobutyrate, iso eugenol acetate, helional, iso E super, ionone gamma methyl, pentalide, galaxolide, phenoxy ethyl propionate.

Other volatile "top note" perfume raw materials having a boiling point of less than, or equal to, 250°C, which should be used in low levels in the present invention, but which have a high odour detection threshold, are selected from, but are not limited to, benzaldehyde, benzyl acetate, camphor, carvone, borneol, bornyl acetate, decyl alcohol, eucalyptol, linalool, hexyl acetate, iso-amyl acetate, thymol, carvacrol, limonene, menthol, iso-amyl alcohol, phenyl ethyl alcohol, alpha pinene, alpha terpineol, citronellol, alpha thujone, benzyl alcohol, beta gamma hexenol, dimethyl benzyl carbinol, phenyl

ethyl dimethyl carbinol, adoxal, allyl cyclohexane propionate, beta pinene, citral, citronellyl acetate, citronellal nitrile, dihydro myrcenol, geraniol, geranyl acetate, geranyl nitrile, hydroquinone dimethyl ether, hydroxycitronellal, linalyl acetate, phenyl acetaldehyde dimethyl acetal, phenyl propyl alcohol, prenyl acetate, triplal, tetrahydrolinalool, verdox, cis-3-hexenyl acetate.

Entrapment Material

Compositions of the present invention comprise an entrapment material at a level such that the weight ratio of the entrapment material to the high odour impact perfume raw materials is in the range of from about 1:20 to about 20:1. It is preferred that the compositions comprise from about 0.1% to about 95%, preferably from about 0.5% to about 50%, more preferably from about 1% to about 25% and even more preferably from about 2% to about 8%, by weight, of an entrapment material.

As defined herein an "entrapment material" is any material which, after application of the composition to a substrate, suppresses the volatility of the perfume raw materials within the fragrance oil thus delaying their evaporation. It is not necessary that the entrapment material forms an association with the perfume raw material within the composition itself, only that this association exists on the substrate after application of the composition. Non limiting examples of mechanisms by which the delay in evaporation may occur are by the entrapment material reversibly or irreversibly, physically or chemically associating with the perfume raw material through complexing, encapsulating, occluding, absorbing, binding, or otherwise adsorbing the perfume raw materials of the fragrance oil.

As defined herein "reversible entrapment" means that any entrapment material: perfume raw material association in which the association can be broken down so that the entrapment material and perfume raw materials are released from each other. As defined herein "irreversible entrapment" means that the entrapment material: perfume raw material association can not be broken down. As defined herein "chemically associated" means that the entrapment material and perfume raw material are linked through a covalent, ionic, hydrogen or other type of chemical bond. As defined herein "physically

"associated" means that the entrapment material and perfume raw material are linked through a bond with a weaker force such as a Van der Waals force. Highly preferred is that, upon the substrate, the entrapment material and the perfume raw material form a reversible physical or chemical association.

As defined herein "to delay the evaporation of a perfume raw material" means to slow down or inhibit the evaporation rate of said perfume raw material from the substrate such that the distinctive fragrance character of the perfume raw material is detectable for at least 2 hours after application to the substrate.

Entrapment materials for use herein are selected from polymers; capsules, microcapsules and nanocapsules; liposomes; film formers; absorbents; cyclic oligosaccharides and mixtures thereof. Preferred are absorbents and cyclic oligosaccharides and mixtures thereof. Highly preferred are cyclic oligosaccharides.

Furthermore it is preferred that, on the substrate, where the perfume raw material and the entrapment material exist in an associated form, that the weight ratio of high odour detection perfume raw material to entrapment material within the associated form is in the range from about 1:20 to about 20:1, preferably from about 1:10 to about 10:1, more preferably in the range from about 1:7 to about 7:1, even more preferably in the range from about 1:5 to about 5:1, further more preferably from about 1:3 to about 3:1 and most preferably in the range from about 1:2 to about 2:1.

Complexation using for example Cyclic Oligosaccharides

It is highly preferred for compositions of the present invention that the entrapment material reversibly, chemically and physically complexes the perfume raw materials. Non limiting, and preferred, examples of entrapment materials that can act in this way are cyclic oligosaccharides, or mixtures of different cyclic oligosaccharides.

As used herein, the term "cyclic oligosaccharide" means a cyclic structure comprising six or more saccharide units. Preferred for use herein are cyclic oligosaccharides having six,

seven or eight saccharide units and mixtures thereof, more preferably six or seven saccharide units and even more preferably seven saccharide units. It is common in the art to abbreviate six, seven and eight membered cyclic oligosaccharides to α , β and γ respectively.

The cyclic oligosaccharide of the compositions used for the present invention may comprise any suitable saccharide or mixtures of saccharides. Examples of suitable saccharides include, but are not limited to, glucose, fructose, mannose, galactose, maltose and mixtures thereof. However, preferred for use herein are cyclic oligosaccharides of glucose. The preferred cyclic oligosaccharides for use herein are α -cyclodextrins or β -cyclodextrins, or mixtures thereof, and the most preferred cyclic oligosaccharides for use herein are β -cyclodextrins.

The cyclic oligosaccharide, or mixture of cyclic oligosaccharides, for use herein may be substituted by any suitable substituent or mixture of substituents. Herein the use of the term "mixture of substituents" means that two or more different suitable substituents can be substituted onto one cyclic oligosaccharide. The derivatives of cyclodextrins consist mainly of molecules wherein some of the OH groups have been substituted. Suitable substituents include, but are not limited to, alkyl groups; hydroxyalkyl groups; dihydroxyalkyl groups; (hydroxyalkyl)alkylenyl bridging groups such as cyclodextrin glycerol ethers; aryl groups; maltosyl groups; allyl groups; benzyl groups; alkanoyl groups; cationic cyclodextrins such as those containing 2-hydroxy-3-(dimethylamino) propyl ether; quaternary ammonium groups; anionic cyclodextrins such as carboxyalkyl groups, sulphobutylether groups, sulphate groups, and succinylates; amphoteric cyclodextrins; and mixtures thereof. Other cyclodextrin derivatives are disclosed in copending application US 09/32192 (27th May 1999), all of which are incorporated herein by reference.

The substituents may be saturated or unsaturated, straight or branched chain. Preferred substituents include saturated and straight chain alkyl groups, hydroxyalkyl groups and mixtures thereof. Preferred alkyl and hydroxyalkyl substituents are selected from C₁-C₈

alkyl or hydroxyalkyl groups or mixtures thereof, more preferred alkyl and hydroxyalkyl substituents are selected from C₁-C₆ alkyl or hydroxyalkyl groups or mixtures thereof, even more preferred alkyl and hydroxyalkyl substituents are selected from C₁-C₄ alkyl or hydroxyalkyl groups and mixtures thereof. Especially preferred alkyl and hydroxyalkyl substituents are propyl, ethyl and methyl, more especially hydroxypropyl and methyl and even more preferably methyl.

Preferred cyclic oligosaccharides for use in the present invention are unsubstituted, or are substituted by only saturated straight chain alkyl, or hydroxyalkyl substituents. Therefore, preferred examples of cyclic oligosaccharides for use herein are α -cyclodextrin, β -cyclodextrin, methyl- α -cyclodextrin, methyl- β -cyclodextrin, hydroxypropyl- α -cyclodextrin and hydroxypropyl- β -cyclodextrin. Most preferred examples of cyclic oligosaccharides for use herein are methyl- α -cyclodextrin and methyl- β -cyclodextrin. These are available from Wacker-Chemie GmbH Hanns-Seidel-Platz 4, Munchen, DE under the tradename Alpha W6 M and Beta W7 M respectively. Especially preferred is methyl- β -cyclodextrin.

Methods of modifying cyclic oligosaccharides are well known in the art. For example, see "*Methods of Selective Modifications of Cyclodextrins*" *Chemical Reviews* (1998) Vol. 98, No. 5, pp 1977-1996, Khan et al and US-A-5, 710, 268.

In addition to preferred substituents themselves, it is also preferred that the cyclic oligosaccharides of the compositions used for the present invention have an average degree of substitution of at least 1.6, wherein the term "degree of substitution" means the average number of substituents per saccharide unit. Preferred cyclic oligosaccharides for use herein have an average degree of substitution of less than about 2.8. More preferably the cyclic oligosaccharides for use herein have an average degree of substitution of from about 1.7 to about 2.0. The average number of substituents can be determined using common Nuclear Magnetic Resonance techniques known in the art.

The cyclic oligosaccharides of the compositions used for the present invention are preferably soluble in both water and ethanol. As used herein "soluble" means at least about 0.1g of solute dissolves in 100ml of solvent, at 25°C and 1 atm of pressure. Preferably the cyclic oligosaccharides for use herein have a solubility of at least about 1g/100ml, at 25°C and 1atm of pressure. Preferred is that cyclic oligosaccharides are only present at levels up to their solubility limits in a given composition at room temperature. A person skilled in the art will recognise that the levels of cyclic oligosaccharides used in the present invention will also be dependent on the components of the composition and their levels, for example the solvents used or the exact fragrance oils, or combination of fragrance oils, present in the composition. Therefore, although the limits stated for the entrapment material are preferred, they are not exhaustive.

Encapsulation using Capsules, Micro-capsules and Nanocapsules

Encapsulation of fragrances within capsules, micro-capsules or nanocapsules which are broken down by environmental triggers can be used to reduce the volatility of fragrance oils by surrounding the oil by small droplets as a resistant wall. This may be either water sensitive or insensitive. In the first case the fragrance can be released when the encapsulated particle is affected by moisture loss from the skin; while in the second case the capsule wall must be ruptured mechanically before the fragrance is released. Encapsulation techniques are well known in the art including DE 1,268,316; US 3,539,465; US 3,455,838.

Moisture sensitive capsules, micro-capsules and nanocapsules are preferably formed from, but not limited to, a polysaccharide polymer. Examples of suitable polymers are dextrans, especially low-viscosity dextrans including maltodextrins. A particularly preferred example of a low viscosity dextrin is one which, as a 50% dispersion in water has a viscosity at 25°C, using a Brookfield Viscometer fitted with an "A" type T-Bar rotating at 20rpm in helical mode, of 330 ± 20 mPa.s. This dextrin is known as Encapsul 855 and is available from National Starch and Chemicals Ltd. A further example of a polysaccharide that can be used to form the moisture sensitive capsules is gum acacia.

Time release micro-capsules are also suitable for use in compositions of the present invention for entrapping hydrophobic perfume raw materials. Such compositions comprise the perfume raw materials encapsulated in a wax or polymer matrix which in turn is coated with a compatible surfactant. The wax or polymers used to form the matrix have a melting point in the range from about 35°C to about 120°C at 1 atmosphere pressure. These are described in detail in EP-A-908,174.

Occlusion using Film Formers

Film formers can also be used to reduce the volatility profile of perfume raw materials. When the fragrance is applied to a substrate, such as the skin, it is believed that film formers entrap the fragrance oils during the evaporation of the volatile solvent thus hindering the release of the volatile material. Any film former which is compatible with the perfume raw materials may be used, preferably the film former will be soluble in water - ethanol mixture. Film former materials useful in this invention include, but are not limited to, ionic and non-ionic derivatives of water soluble polymers. Examples of suitable film forming materials are water soluble polymers containing a cationic moiety such as polyvinyl pyrrolidine and its derivatives having a molecular weight of 50,000 to 1,000,000. Other examples of ionic polymeric film forming materials are cationic cellulose derivatives sold under the trade names of Polymer JR (union Carbide), Klucel GM (hercules) and ethoxylated polyethyleneimine sold under the trade name PEI 600 (Dow). Examples of suitable cellulosic derivatives such as hydroxymethyl cellulose, hydroxypropyl methylcellulose and hydroxyethyl cellulose. Another examples of film formers is benzophenone.

Non limiting examples of film forming materials are given in US 3,939,099.

Other Polymers

Additional non limiting examples of other polymer systems that can be used include water soluble anionic polymers eg polyacrylic acids and their water-soluble salts are useful in the present invention to delay the evaporation rate of certain amine-type odours.

Preferred polyacrylic acids and their alkali metal salts have an average molecular weight

of less than about 20,000, preferably less than 10,000, more preferably from about 500 to about 5,000. Polymers containing sulphonic acid groups, phosphoric acid groups, phosphonic acid groups and their water soluble salts, and their mixtures thereof, and mixtures with carboxylic acid and carboxylate groups, are also suitable.

Water soluble polymers containing both cationic and anionic functionalities are also suitable. Examples of these polymers are given in US Patent 4,909,986. Another example of water-soluble polymers containing both cationic and anionic functionalities is a copolymer of dimethyldiallyl ammonium chloride and acrylic acid, commercially available under the trade name Merquat 280® from Calgon.

Absorption of Perfume Raw Material

When clarity of solution is not needed, odour absorbing materials such as zeolites and / or activated carbon can be used to modify the release rate of perfume raw materials.

A preferred class of zeolites is characterised as "intermediate" silicate / aluminate zeolites. The intermediate zeolites are characterised by $\text{SiO}_2 / \text{AlO}_2$ molar ratios of less than about 10, preferably in the range from about 2 to about 10. The intermediate zeolites have an advantage over the "high" zeolites since they have an affinity for amine-type odours, they are more weight efficient for odour absorption since they have a larger surface area and they are more moisture tolerant and retain more of their odour absorbing capacity in water than the high zeolites. A wide variety of intermediate zeolites suitable for use herein are commercially available as Valfor® CP301-68, Valfor® 300-63, Valfor® CP300-35 and Valfor® 300-56 available from PQ Corporation, and the CBV100® series of zeolites from Conteka. Zeolite materials marketed under the trade name Abscents® and Smellrite® available from The Union Carbide Corporation and UOP are also preferred. These materials are typically available as a white powder in the 3-5 μm particle size range. Such materials are preferred over the intermediate zeolites for control of sulphur containing odours eg thiols, mercaptans.

Carbon materials suitable for use in the present invention are materials well known in commercial practice as absorbents for organic molecules and / or for air purification purposes. Often, such carbon material is referred to as "activated" carbon or "activated charcoal". Such carbon is available from commercial sources under trade names as; Calgon-Type CPG®; Type PCB®; Type SGL®; Type CAL®; and Type OL®.

Other odour absorbers suitable for use herein include silica molecular sieves, activated alumina, bentonite and kaolonite.

Volatile Solvent

It is preferred that compositions of the present invention comprise greater than about 50%, preferably from about 55% to about 99.9%, more preferably from about 60% to about 95%, even more preferably from about 65% to about 75%, by weight, of a volatile solvent or mixture of volatile solvents. Any volatile solvent suitable for use in the compositions can be used herein. The solvents for use herein are preferably organic volatile solvents.

As used herein, "volatile" refers to substances with a significant amount of vapour pressure under ambient conditions, as is understood by those in the art. The volatile solvents for use herein will preferably have a vapour pressure of about 2kPa or more, more preferably about 6kPa or more at 25°C. The volatile solvents for use herein will preferably have a boiling point under 1 atm, of less than about 150°C, more preferably less than about 100°C, even more preferably less than about 90°C, even more preferably still less than about 80°C.

Preferably the volatile solvents for use herein will be safe for use on a wide range of substrates, more preferably on human or animal skin or hair. Suitable volatile solvents include, but are not limited to, those found in the *CTFA International Cosmetic Ingredient Dictionary and Handbook, 7th edition, volume 2 P1670 - 1672, edited by Wenninger and McEwen (The Cosmetic, Toiletry, and Fragrance Association, Inc., Washington, D. C., 1997)*. Conventionally used volatile solvents include C₃-C₁₄ saturated

and unsaturated, straight or branched chain hydrocarbons such as cyclohexane, hexane, heptane, isoctane, isopentane, pentane, toluene, xylene; halogenated alkanes such as perfluorodecalin; ethers such as dimethyl ether, diethyl ether; straight or branched chain alcohols and diols such as methanol, ethanol, propanol, isopropanol, n-butyl alcohol, t-butyl alcohol, benzyl alcohol, butoxypropanol, butylene glycol, isopentyldiol; aldehydes and ketones such as acetone; volatile silicones such as cyclomethicones for example octamethyl cyclo tetrasiloxane and decamethyl cyclopentane siloxane; volatile siloxanes such as phenyl pentamethyl disiloxane, phenylethylpentamethyl disiloxane, hexamethyl disiloxane, methoxy propylheptamethyl cyclotetrasiloxane, chloropropyl pentamethyl disiloxane, hydroxypropyl pentamethyl disiloxane, octamethyl cyclotetrasiloxane, decamethyl cyclopentasiloxane; propellants, and mixtures thereof. Preferred volatile solvents are ethers such as dimethyl ether, diethyl ether; straight or branched chain alcohols and diols such as methanol, ethanol, propanol, isopropanol, n-butyl alcohol, t-butyl alcohol, benzyl alcohol, butoxypropanol, butylene glycol, isopentyldiol; volatile silicones such as cyclomethicones for example octamethyl cyclo tetrasiloxane and decamethyl cyclopentane siloxane; propellants, and mixtures thereof. More preferred for use herein are C₁-C₄ straight chain or branched chain alcohols for example methanol, ethanol, propanol, isopropanol and butanol and mixtures thereof, and most preferred for use herein is ethanol.

Balance of the Composition

The compositions herein are useful for use a fragrance component to a wide variety of different types of compositions including those such as cosmetic compositions, fragrance compositions for human or animal use, air freshener compositions, aromatherapy compositions, laundry and cleaning compositions, fragranced paper, cloth or other substrate type products and the like, in fact in any type of composition which comprises a fragrance. In all of these cases the compositions would comprise a variety of other optional ingredients which would render the compositions more acceptable or provide them with additional usage benefits. Such conventional optional ingredients are well-known to those skilled in the art and will vary greatly depending upon the field in which the compositions are to be used. Although compositions according to the second aspect

of this invention comprise greater than 50% volatile solvent they should still be considered as being useful for inclusion in a wide variety of compositions with both cosmetic and non cosmetic benefits.

For cosmetic compositions in particular, the compositions can comprise any cosmetically acceptable ingredients such as those found in the *CTFA International Cosmetic Ingredient Dictionary and Handbook, 7th edition, edited by Wenninger and McEwen, (The Cosmetic, Toiletry, and Fragrance Association, Inc., Washington, D.C., 1997)*. As used herein "cosmetically acceptable" means a material (e.g., compound or composition) which is suitable for use in contact with skin, hair or other suitable substrate as defined herein below. However that these ingredients are listed as useful for inclusion within a cosmetic composition does not preclude them from being incorporated into any other type of composition which comprises a fragrance formulated according to the present invention.

Pro-Perfumes

Compositions of the present invention optionally comprise pro-perfumes, or pro-fragrances. Synthesising pro-perfumes or pro-fragrances from perfume raw materials can result in compounds which impart a delayed release mechanism to that specific perfume raw material. Pro-perfumes or pro-fragrances are not considered as an entrapment material within this invention and if added to the composition should be considered as a further additional ingredient. Pro-perfumes useful within the present invention include those selected from more than 1 type of pro-chemistry which ensures that a wide range of possible perfume raw materials can be used. This is consistent with the objective of providing unique fragrances with a broad spectrum of "top note" characters.

Within a pro-perfume the perfume raw material has been reacted with more than one type of chemical groups such as acetal, ketal, ester, hydrolysable inorganic -organic. As such, as defined within the present invention, the perfume raw material is considered to constitute part of the fragrance oil and the chemical groups to constitute part of the entrapment material. Pro-perfumes themselves are designed to be non-volatile, or else have a very low volatility. However, once on the substrate, the perfume raw material is

released from the pro-perfume. Once released the perfume raw material has its original characteristics. The perfume raw material may be released from the pro-perfume in a number of ways. For example, it may be released as a result of simple hydrolysis, or by shift in an equilibrium reaction or by a pH-change, or by enzymatic release. The fragrances herein can be relatively simple in their compositions, comprising a single chemical, or can comprise highly sophisticated complex mixtures of natural and synthetic chemical components, all chosen to provide any desired odour.

Non-limiting pro-perfumes suitable for use in the present application are described in WO 98/47477, WO 99/43667, WO 98/07405, WO 98/47478 and copending applications US 60/105380 (23rd Oct 1998) and US 60/130108 (20th April 1999).

Nonvolatile solvents

While the compositions of the present invention must comprise a volatile solvent they may also comprise "nonvolatile" solvents. Suitable non-volatile solvents include, but are not limited to, benzyl benzoate, diethyl phthalate, isopropyl myristate, and mixtures thereof.

Molecular wedges

When cyclic oligosaccharides are present in the compositions of the present invention, low molecular weight polyol molecular wedge having from about 2 to about 12 carbon atoms, preferably from about 2 to about 6 carbon atoms and at least one -OH functional group, preferably at least 2 -OH functional groups are preferably used herein for further prolonging the fragrance character of the composition. These polyols can further contain ether groups within the carbon chain. Suitable examples include ethylene glycol, propylene glycol, dipropylene glycol, 1,4-butanediol, 1,6-hexanediol and mixtures thereof. When present these polyols are present at a level of from about 0.01% to about 20%, preferably from about 0.1% to about 10%, and especially from about 0.5% to about 5% by weight of composition. It is preferred that the molar ratio of molecular wedge material to oligosaccharide is from 10:1 to 1:10, preferably 1:1 or greater, especially 1:1.

While not wishing to be limited by theory, the above mentioned molecular wedge molecules can form tertiary inclusion complexes with the complexed perfume material and the cyclic oligosaccharide. These small dipolar molecules can fit into the cavity of the cyclic oligosaccharide and anchor via their OH groups onto the outside rim of the cyclic oligosaccharide through hydrogen bonding. This enables the inclusion of all or parts of the fragrance material into the cavity of the cyclic oligosaccharide such that the stability of the formed tertiary complex is increased versus the complex formed by the fragrance material and cyclic oligosaccharide alone.

Water

The compositions of the present invention may also comprise water. If present, the water will preferably comprise from about 0.1% to about 40%, more preferably from about 1% to about 30%, even more preferably about 5% to about 20%, by weight, of total composition. When the composition is a cosmetic composition the level of water should not be so high that the product becomes cloudy thus negatively impacting the product aesthetics.

There are a number of other examples of additional ingredients that are suitable for inclusion in the present compositions, particularly in compositions for cosmetic use. These include, but are not limited to, alcohol denaturants such as denatonium benzoate; UV stabilisers such as benzophenone-2; antioxidants such as tocopheryl acetate; preservatives such as phenoxyethanol, benzyl alcohol, methyl paraben, propyl paraben; dyes; pH adjusting agents such as lactic acid, citric acid, sodium citrate, succinic acid, phosphoric acid, sodium hydroxide, sodium carbonate; deodorants and anti-microbials such as farnesol and zinc phenolsulphonate; humectants such as glycerine; oils; skin conditioning agents such as allantoin; cooling agents such as trimethyl isopropyl butanamide and menthol; hair conditioning ingredients such as panthenol, panthetine, pantotheine, panthenyl ethyl ether, and combinations thereof; silicones; solvents such as hexylene glycol; hair-hold polymers such as those described in WO-A-94/08557; salts in general, such as potassium acetate and sodium chloride and mixtures thereof. If present, these additional ingredients will preferably be present at a level of less than 10%, by

weight, of total composition. More preferably these additional ingredients will be present at a level of less than about 5%, by weight, of total composition.

Product Forms

The compositions for use in the present invention may take any form suitable for use, more preferably for cosmetic use. These include, but are not limited to, vapour sprays, aerosols, emulsions, lotions, liquids, creams, gels, sticks, ointments, pastes, mousses, powders, granular products, substrates, cosmetics (eg semi-solid or liquid make-up, including foundations) and the like. Preferably the compositions for use in the present invention take the form of a vapour spray. Compositions of the present invention can be further added as an ingredient to other compositions, preferably cosmetic compositions, in which they are compatible. As such they can be used within solid composition or applied substrates etc.

The compositions for use in the present invention will preferably comprise an acceptable carrier, preferably a cosmetically acceptable carrier. The phrase "carrier", as used herein, means one or more compatible solid or liquid fillers, diluents, extenders and the like. The term "compatible", as used herein, means that the components of the compositions of this invention are capable of being combined with the primary actives of the present invention, and with each other, in a manner such that there is no interaction which would substantially reduce the efficacy of the composition under ordinary use situations. The type of carrier utilised in the present invention depends of the type of product desired and may comprise, but are not limited to, solutions, aerosols, emulsions (including oil-in-water or water-in-oil), gels, solids and liposomes.

Preparation of Composition

Compositions and fragrance oils for use in the present invention should be prepared according to procedures usually used and that are well known and understood by those skilled in the art with materials of similar phase partitioning can be added in any order. The entrapment of the perfume raw materials can occur at any reasonable stage in the preparation of the overall composition. As such the fragrance oil can be prepared in its

entirety, then entrapped with a suitable material before addition to the remainder of the composition. Alternatively the entrapment material can be added to the balance of the composition prior to addition of the complete fragrance oil. Finally it is possible to entrap any single perfume raw material, or group of raw materials, individually before either adding these to the balance of the fragrance oil or to the balance of the composition.

Methods of Use

The present invention preferably relates to compositions, particularly cosmetic compositions, and more particularly fragrance compositions, which comprise a fragrance oil, in combination with an entrapment material, such that the fragrance profile preferably remains detectable for at least about 2 hours after applying the composition to the substrate.

Therefore according to another aspect of the present invention there is provided a composition comprising:

- (a) a fragrance oil wherein the fragrance oil comprises:
 - (i) one or more perfume raw materials with a high odour impact which have an odour detection threshold of less than, or equal to about 50 parts per billion
 - (ii) less than about 5%, by weight of the fragrance oil, of top note perfume raw materials where in the top note perfume raw materials have a boiling point of less than 250°C at 1 atmosphere pressure
- (b) an entrapment material which is selected from the group consisting of polymers; capsules, microcapsules, and nanocapsules; liposomes; film formers; absorbents; cyclic oligosaccharides and mixtures thereof;

wherein the perfume raw material and the entrapment material exist in an associated form on the substrate, wherein the weight ratio of high odour impact perfume raw materials which have an odour detection threshold of less than, or equal to, about 50 parts per billion to entrapment material within the associated form falls in the range of about 1:20 to about 20:1 and wherein the high odour impact raw material, or mixture of raw materials, remain detectable on the substrate for greater than about 2 hours, preferably

greater than about 4 hours, more preferably greater than about 6 hours after the composition has been applied to the substrate.

The composition itself is preferably used for providing fragrance to a suitable substrate. As used herein the term "suitable substrate" means any surface to which the present composition may be applied without an unduly adverse effect. This can include a wide range of substrates including human or animal skin or hair, paper, furnishings, materials, and can also include the air in a room. Preferred substrates are the skin or hair, especially the skin.

The compositions of the present invention may be used in a conventional manner for fragrancing a suitable substrate. An effective amount of the composition, typically from about 1 μ l to about 1000 μ l, preferably from about 10 μ l to about 250 μ l, more preferably from about 25 μ l to about 100 μ l, is applied to the substrate. The composition may be applied by hand but is preferably applied utilising a vaporiser. Preferably, the composition is then left to dry. The scope of the present invention should be considered to cover one or more distinct applications of a product or the continuous release of a composition via a vaporiser or other type of atomiser.

Examples

The following examples further illustrate the preferred embodiments within the scope of the present invention. These examples are given solely for the purposes of illustration and are not to be construed as limitations of the present invention as many variations of the invention are possible without departing from its spirit or scope. Unless otherwise indicated, all ingredients are expressed on a weight percentage of the active ingredient.

Fragrance Oil Examples I-V

Examples I-V are non limiting examples of the fragrance oil.

<u>Perfume Raw Material</u>	<u>I (%)</u>	<u>II(%)</u>	<u>III(%)</u>	<u>IV(%)</u>	<u>V(%)</u>
Methyl Cedrylone ¹	5.00	0.00	8.35	15.00	15.00
Benzoin Claire 50% in dep ²	5.00	2.00	10.00	5.00	2.00
Benzyl Salicylate ³	20.00	10.00	10.00	10.00	18.00
Benzyl Benzoate ⁴	20.00	10.00	10.00	10.00	10.00
Methyl dihydro Jasmonate ⁵	25.00	40.00	20.00	14.35	14.10
Cis 3- Hexenyl Salicylate ¹	5.00	5.00	5.00	5.00	5.00
Aurantiol ¹	3.00	3.00	3.00	3.00	1.00
Ethyl Vanillin 10% dpg ⁶	0.10	0.10	0.10	0.10	0.10
Cymal ⁷	5.00	5.00	5.00	5.00	5.00
Vanillin ⁶	1.00	1.00	6.00	3.00	2.00
Ionone Beta ⁸	1.00	0.25	0.25	0.25	2.00
Ethyl Maltol 10 % dpg ⁹	0.10	0.10	2.00	0.10	0.10
Eugenol ¹⁰	0.10	0.10	0.10	4.00	0.50
Cinnamic Alcohol ⁷	2.00	2.00	2.00	2.00	2.00
Methyl	0.20	0.20	0.20	0.20	0.20

Cinnamate ⁴					
Ethyl Methyl Phenyl Glycidate ¹¹	0.50	5.00	3.00	3.00	3.00
Pentalide ¹	5.00	15.00	15.00	15.00	15.00
Ethylene Brassylate ¹²	2.00	1.25	0.00	0.00	0.00
Cedrol ¹	-	-	-	5.00	5.00
	100.00	100.00	100.00	100.00	100.00

1 Fragrance Resources Inc. 620 Route 3 West PO Box 4277 Clifton, NJ 07012

2 Robertet Inc. 125 Bauer Drive Oakland, NJ 07436

3 Rhone Poulenc Inc. CN 5266 Princeton , NJ 08543-5266

4 H&R 300 North Street Teterboro, NJ 07608

5 Firmenich SA, 1 Route des Jeunes, CH-1211 Geneva 8 SWITZERLAND

6 Polarome 200 Theodore Conrad Drive, Jersey City, NJ 07305

7 Givaudan 19-23 voie des Bans BP98, 95101 Argenteuil Cedex, FRANCE

8 International Flavors & Fragrances 521 W. 57th St, New York, NY 10019 USA

9 Pfizer 9 Pfizer - Danisco Coltor 430 Saw Mill River Rd. Ardsley, NY 10502

10 Manheimer 47-22 Pearson Place , Long Island Cirt, NY 11101

11 Dragoco Gerberding & Co AG, D-37601 Holzminden GERMANY

12 Takasgo International 4 Volvo Drive PO Box 932 Rockleigh NJ, 07647

The perfume raw materials were mixed with stirring at room temperature.

Alcoholic Compositions Examples VI-X

	VI (%)	VII (%)	VIII (%)	IX (%)	X (%)
Fragrance (selected from examples I-V)	1.5	5	4	7	12
Cyclic	2.5	5	4	5	6

Oligosaccharide ¹³					
Ethanol	to 100				
Deionised Water	15.8	12.9	13.7	12.9	-

¹³ Beta W7 M available from Wacker-Chemie GmbH, Hanns-Seidel-Platz 4, Munchen, DE

The cyclic oligosaccharide was dissolved in ethanol at room temperature with stirring.
Then the fragrance and water were added with stirring.

Alcoholic Compositions Examples XI-XV

	XI (%)	XII (%)	XIII (%)	XIV (%)	XV (%)
Fragrance (selected from examples I-V)	1.5	5	4	7	12
Polyoxymethylene Urea (PMU) microcapsules ¹⁴	2.0	3	2	5	6
Ethanol	to 100	to 100	to 100	to 100	to 100
Deionised Water	15.8	12.9	13.7	12.9	12.8

¹⁴ Polyoxymethylene Urea (PMU) microcapsules available from 3M Center, Building 275-5E-08, St. Paul,
MN 55144-1000, USA

Cosmetic Compositions Examples XVI-XVIII

	XVI (%)	XVII (%)	XVIII (%)
Fragrance (selected from examples I-V)	3.2	2	3
Cyclic Oligosaccharide ¹³	2.5	1.5	3.5
Zinc phenolsulphonate	2	1	2
Dipropylene Glycol	30.5	19	14
Isopropyl myristate	1.5	6	7
Ethanol	to 100	to 100	to 100

¹³ Beta W7 M available from Wacker-Chemie GmbH, Hans-Seidel-Platz 4, Munchen, DE

The zinc phenolsulphonate is stirred into the ethanol until fully dissolved. Then the dipropylene glycol is added with stirring. Next the isopropyl myristate, then the cyclic oligosaccharide and then the fragrance are all added with stirring. For an aerosol deodorant a propellant such as propane butane (CAP 40®) can be added to Examples XVI-XVIII according to standard industry practice.

When examples VI-XVIII were applied to the substrate the fragrance characters could still be determined for three to six hours, preferably longer, after application. By contrast, the same long lasting effect was not achieved when control compositions, comprising the same fragrance oil but without the entrapment material, were applied to a substrate.

Claims:

1. A composition comprising:
 - (a) a fragrance oil wherein the fragrance oil comprises:
 - (i) one or more perfume raw materials with a high odour impact which have an odour detection threshold of less than, or equal to about 50 parts per billion
 - (ii) less than about 5%, by weight of the fragrance oil, of top note perfume raw materials where in the top note perfume raw materials have a boiling point of less than 250°C at 1 atmosphere pressure
 - (b) an entrapment material which is selected from the group consisting of polymers; capsules, microcapsules, and nanocapsules; liposomes; film formers; absorbents; cyclic oligosaccharides and mixtures thereof;
wherein the perfume raw material and the entrapment material exist in an associated form on the substrate and wherein the weight ratio of high odour impact perfume raw materials which have an odour detection threshold of less than, or equal to, about 50 parts per billion to entrapment material within the associated form falls in the range of about 1:20 to about 20:1.
2. A composition according to Claim 1 wherein the weight ratio of the high odour impact perfume raw materials to the entrapment material within the associated form is in the range of from about 1:10 to about 10:1, preferably from about 1:7 to about 7:1, more preferably from about 1:5 to about 5:1, even more preferably from about 1:3 to about 3:1 and most preferably from about 1:2 to about 2:1.
3. A composition according to Claim 1 or 2 wherein the composition comprises from about 0.01% to about 99%, preferably from about 0.25% to about 50%, more preferably from about 0.5% to about 40%, even more preferably from about 1% to about 25%, and most preferably from about 2.5% to about 25%, by weight, of the fragrance oil.
4. A composition according to Claim 1 or 3 wherein the fragrance oil comprises from about 0.1% to about 99%, preferably from about 1% to about 75%, more preferably from about 5% to about 50%, and even ore preferably from about 20% to about 40%,

by weight of the fragrance oil, of high odour impact perfume raw materials which have an odour detection threshold of less than, or equal to, 50 parts per billion.

5. A composition according to any of Claims 1 to 4 wherein the high odour impact perfume raw materials have an odour detection threshold of less than, or equal to about 40 parts per billion, preferably about 30 parts per billion, more preferably about 20 parts per billion and even more preferably about 10 parts per billion.
6. A composition according to any of Claims 1 to 4 wherein the fragrance oil comprises less than about 4%, preferably less than about 3% and more preferably less than about 2%, by weight of fragrance oil, of top note perfume raw materials.
7. A composition according to any of Claims 1 to 6 wherein the entrapment material is selected from the group consisting of absorbents; cyclic oligosaccharides and mixtures thereof.
8. A composition according to any of Claims 1 to 7 wherein the entrapment material is selected from cyclic oligosaccharides.
9. A composition according to any of Claims 1 to 8 wherein the cyclic oligosaccharide has six, seven or eight saccharide units and mixtures of thereof, preferably six or seven saccharide units and more preferably seven saccharide units.
10. A composition according to any of Claims 1 to 9 wherein the cyclic oligosaccharide is a cyclodextrin.
11. A composition according to any of Claims 1 to 10 wherein the cyclic oligosaccharide is substituted with a substituent selected from C₁-C₈ alkyl or hydroxyalkyl groups and mixtures thereof, preferably C₁-C₄ alkyl or hydroxyalkyl groups and mixtures thereof.
12. A composition according to any of Claims 1 to 11 wherein the cyclic oligosaccharide is preferably substituted with C₁-C₄ alkyl groups and more preferably methyl substituted.
13. A composition according to any of Claims 1 to 12 wherein the cyclic oligosaccharide is α-cyclodextrin, β-cyclodextrin, methyl-α-cyclodextrin, methyl-β-cyclodextrin, or mixtures thereof.
14. A composition according to any of Claims 1 to 13 wherein the cyclic oligosaccharide is methyl-β-cyclodextrin.

15. A composition according to any of Claims 1 to 14 which comprises from about 0.1% to about 95%, more preferably from about 0.5% to about 50%, even more preferably from about 1% to about 25%, and most preferably from about 2% to about 8%, by weight, of said entrapment material.
16. A composition according to any of Claims 1 to 15 wherein the volatile solvent has a boiling point, at 1 atmosphere pressure, of less than about 150°C, more preferably less than about 100°C, even more preferably less than about 90°C, even more preferably still less than about 80°C.
17. A composition according to any of Claims 1 to 16 wherein the volatile solvent is selected from ethers; straight or branched chain alcohols and diols; volatile silicones; propellants, and mixtures thereof, preferably C₁-C₄ straight chain or branched chain alcohols and mixtures thereof.
18. A composition according to any of Claims 1 to 17 wherein the volatile solvent is ethanol.
19. A composition according to any of Claims 1 to 18 which comprises greater than about 50%, preferably from about 55% to about 99.9%, more preferably from about 60% to about 95%, even more preferably from about 65% to about 75%, by weight, of volatile solvent.
20. A composition according to any of Claims 1 to 19 wherein the composition is a cosmetic composition, preferably a fine fragrance composition.
21. A composition comprising:
 - (a) a fragrance oil wherein the fragrance oil comprises:
 - (i) one or more perfume raw materials with a high odour impact which have an odour detection threshold of less than, or equal to about 50 parts per billion
 - (ii) less than about 5%, by weight of the fragrance oil, of top note perfume raw materials where in the top note perfume raw materials have a boiling point of less than 250°C at 1 atmosphere pressure
 - (b) an entrapment material which is selected from the group consisting of polymers; capsules, microcapsules, and nanocapsules; liposomes; film formers; absorbents; cyclic oligosaccharides and mixtures thereof;

wherein the perfume raw material and the entrapment material exist in an associated form on the substrate, wherein the weight ratio of high odour impact perfume raw materials which have an odour detection threshold of less than, or equal to, about 50 parts per billion to entrapment material within the associated form falls in the range of about 1:20 to about 20:1 and wherein the high odour impact raw material, or mixture of raw materials, remain detectable on the substrate for greater than about 2 hours, preferably greater than about 4 hours, more preferably greater than about 6 hours after the composition has been applied to the substrate.

22. A method of fragrancing a substrate comprising:

- (i) applying a composition according to any of Claims 1 to 21 to said substrate; and preferably
- (ii) allowing said composition to dry.

23. Use of a composition according to any of Claims 1 to 21 for fragrancing a substrate for at least about 2 hours, preferably at least about 4 hours, more preferably at least about 6 hours.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/I 10/29759

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61K7/46

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

WPI Data, EPO-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 98 56429 A (THE PROCTER & GAMBLE COMPANY) 17 December 1998 (1998-12-17) page 20, line 30 -page 24, line 20 page 45, line 1 -page 50, line 4 examples 19,21,23,25,27,29,31,35,38	1-4, 6-15,20, 22
Y	WO 99 55819 A (THE PROCTER & GAMBLE COMPANY ET AL.) 4 November 1999 (1999-11-04) page 3 -page 9 page 10, paragraph 2 claims 1,2	1-4, 6-15, 21-23
Y	WO 98 56341 A (THE PROCTER & GAMBLE COMPANY) 17 December 1998 (1998-12-17) page 4 page 10, paragraph 4 -page 13, paragraph 4	1-4, 6-15,22
		-/-

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INTERNATIONAL SEARCH REPORT

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 5 578 563 A (TOAN TRINH ET AL.) 26 November 1996 (1996-11-26) column 9, line 40 -column 11, line 14 -----	7-14
Y	PATENT ABSTRACTS OF JAPAN vol. 199, no. 810, 31 August 1998 (1998-08-31) & JP 10 120541 A (KAO CORP. ET AL.), 12 May 1998 (1998-05-12) cited in the application abstract -----	21,23

INTERNATIONAL SEARCH REPORT
in relation to patent family members

International Application No

PCT/I 10/29759

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 9856429	A	17-12-1998	US 5955093 A US 5997759 A US 5942217 A US 6033679 A AU 4356997 A AU 7961098 A BR 9810426 A CN 1269819 T EP 0988064 A EP 0988364 A EP 0988365 A TR 200000439 T US 5968404 A WO 9856337 A WO 9856888 A WO 9856889 A WO 9856890 A US 6001343 A US 6106738 A AU 1711199 A BR 9815837 A WO 9955815 A AU 1711099 A AU 1804699 A BR 9815835 A BR 9815836 A TR 200003126 T WO 9955813 A WO 9955814 A	21-09-1999 07-12-1999 24-08-1999 07-03-2000 30-12-1998 30-12-1998 05-09-2000 11-10-2000 29-03-2000 29-03-2000 29-03-2000 21-11-2000 19-10-1999 17-12-1998 17-12-1998 17-12-1998 17-12-1998 14-12-1999 22-08-2000 16-11-1999 26-12-2000 04-11-1999 16-11-1999 16-11-1999 26-12-2000 26-12-2000 22-01-2001 04-11-1999 04-11-1999
WO 9955819	A	04-11-1999	AU 3048199 A BR 9909803 A EP 1073705 A	16-11-1999 26-12-2000 07-02-2001
WO 9856341	A	17-12-1998	US 5942214 A US 5861147 A AU 7825798 A BR 9809995 A CN 1261785 T EP 0991401 A NO 996055 A TR 9903041 T	24-08-1999 19-01-1999 30-12-1998 01-08-2000 02-08-2000 12-04-2000 04-02-2000 21-06-2000
US 5578563	A	26-11-1996	CA 2197442 A EP 0774979 A JP 10505257 T JP 3090951 B WO 9604938 A	22-02-1996 28-05-1997 26-05-1998 25-09-2000 22-02-1996
JP 10120541	A	12-05-1998	NONE	

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